

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: AUSTIN PHARMA,

LLC

[Docket No. DEA-392]

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated his authority under the Controlled Substances

Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR

0.100(b). Authority to exercise all necessary functions with respect to the promulgation
and implementation of 21 CFR part 1301, incident to the registration of manufacturers,
distributors, and dispensers of controlled substances (other than final orders in connection
with suspension, denial, or revocation of registration) has been redelegated to the Deputy

Assistant Administrator of the DEA Office of Diversion Control ("Deputy Assistant
Administrator") pursuant to sec. 7(g) of 28 CFR pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.33(a), this is notice that on March 27, 2014, Austin Pharma, LLC., 811 Paloma Drive, Suite C, Round Rock, Texas 78665-2402, applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	<u>Schedule</u>
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I

The company plans to manufacture bulk synthetic active pharmaceutical ingredients (APIs) for distribution and new product development to its customers. The company plans to bulk manufacture a synthetic tetrahydrocannabinol.

In reference to drug code 7360, the company plans to manufacture a synthetic cannabinol in bulk for sale to its customers. The controlled substance will be further synthesized to bulk manufacture a synthetic tetrahydrocannabinol (7370). No other activity for this drug code is authorized for this registration.

Dated: May 28, 2014.

Joseph T. Rannazzisi, Deputy Assistant Administrator.

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